



Europäisches Patentamt
European Patent Office
Office européen des brevets

(11) Publication number:

**0 253 410
A1**

(12)

EUROPEAN PATENT APPLICATION

(21) Application number: 87110386.7

(51) Int. Cl.4: **C07C 21/18**, **C07C 17/00**

(22) Date of filing: 17.07.87

(30) Priority: 18.07.86 IT 2117186

(43) Date of publication of application:
20.01.88 Bulletin 88/03

(54) Designated Contracting States:
BE DE ES FR GB IT NL SE

(71) Applicant: **AUSIMONT S.p.A.**
31, Foro Buonaparte
I-20121 Milano(IT)

(72) Inventor: **Gervasutti, Claudio**
2/4, via Paolo Sarpi
I-30175 Mestre-Venezia(IT)

(74) Representative: **Barz, Peter, Dr. et al**
Patentanwälte Dipl.-Ing. G. Dannenberg Dr.
P. Weinhold, Dr. D. Gudel Dipl.-Ing. S.
Schubert, Dr. P. Barz Siegfriedstrasse 8
D-8000 München 40(DE)

(54) **Process for preparing fluoroethylenes and chlorofluoroethylenes from chlorofluoroethanes.**

(57) A process for preparing fluoroethylenes and chlorofluoroethylenes starting from chlorofluoroethanes and hydrogen, carried out in gas phase at a temperature ranging from 150 to 600°C, in the presence of a hydrogenation catalyst, wherein the chlorofluoroethanes have at least one chlorine atom on each carbon atom, and the catalyst is selected preferably from palladium, nickel, chromium, cobalt, platinum, copper and mixtures thereof, and it is utilised either as such or on inert materials.

EP 0 253 410 A1

PROCESS FOR PREPARING FLUORETHYLENES AND CHLOROFLUOROMETHYLENES FROM
CHLOROFLUOROETHANES

This invention relates to a process for preparing fluoroethylenes and chlorofluoroethylenes by hydrogenolysis of chlorofluoroethanes, comprising also the compounds which contain, besides chlorine and fluorine, one or more hydrogen atoms.

Fluoroethylenes and chlorofluoroethylenes are halogenated olefins well known in literature and they are advantageously utilized as intermediates in the preparation of fluoroplastomers, fluoroelastomers and as comonomers in the preparation of fluorinated copolymers.

Fluorine-containing olefins are generally prepared by dehydrohalogenation or dehalogenation of the corresponding haloalkanes in the liquid phase, according to well known procedures. For example, 1-chloro-1,2-difluoroethylene and 1,2-dichloro-dichloroethylene are prepared by reduction, with zinc in an alcohol solution, 1,2-difluoro-1,1,2-trichloroethane and 1,2-difluoro-tetrachloroethane, respectively (J. Chem. Soc. London 1957, pages 2800-06). The high addition of zinc necessary to the reaction and its difficult removal render these processes little attractive from an industrial viewpoint.

The preparation of 1,2-difluoroethylene using 1,2-difluoro-tetrachloroethane as a starting material is described in literature. Said reaction occurs in two steps, the former comprising the obtainment of 1,2-difluoro-1,2-dichloroethane through reduction by means of LiAlH_4 or through UV-radiations in the presence of 2-propanol, while the latter consists in the dehalogenation of the abovesaid intermediate in the presence of Mg (Collection Czechoslov. Chem. Commun. Vol. 39, (1974) pages 2801-07).

It has now surprisingly been found that fluoroethylenes and chlorofluoroethylenes of the above-cited type are preparable in a single step by reaction, in a gas phase, of chlorofluoroethanes with hydrogen, in the presence of hydrogenation catalysts.

Thus, the object of the present invention is a process for preparing fluoroethylenes and chlorofluoroethylenes, comprising reacting hydrogen with chlorofluoroethanes having at least one chlorine atom on each carbon atom, in gas phase, at a temperature ranging from 150 to 600°C and in the presence of hydrogenation catalysts.

The process of the present invention is an improvement as compared with those of the art because of its profitability, easy operability on a commercial scale and because it permits to prepare fluoroethylenes and chlorofluoroethylenes in a single step.

Said process can be carried out continuously and permits to obtain a very high conversion of the reagents.

Utilizable chlorofluoroethanes for the present invention are the ones, as already mentioned, containing at least one chlorine atom on each carbon atom, in particular 1,2-difluorotetrachloroethane, 1,1-difluorotetrachloroethane, 1,1,2-trichlorotrifluoroethane.

Any hydrogenation catalyst can be used for the process of the present invention; examples thereof are palladium, nickel, chromium, cobalt, platinum, copper, which can be used either pure or in admixture. In particular, palladium and nickel are preferred.

Such catalysts can be utilized either as such or, preferably, on inert materials such as, e.g., carbon, aluminium oxide, barium sulphate at a concentration ranging from 0.1 to 10% by weight.

Hydrogenation is carried out in tubular reactors made of materials such as, for example, nickel, Inconel, stainless steel.

The hydrogenolysis temperature ranges from 150°C to 600°C, preferably from 200°C to 400°C. The process can be carried out either at atmospheric pressure or at a higher pressure.

The contact time of the reagents with the catalyst generally ranges from 5 to 60 seconds, preferably from 10 to 20 seconds.

Hydrogen can be fed either in the pure state or diluted with an inert gas such as for example nitrogen, helium, argon.

The molar ratio between hydrogen and chlorofluoroethanes ranges from 1.0 to 10, preferably from 2 to 5, and it is selected as a function of the product to be obtained; the higher said molar ratio is, the more the product mixture gets rich in hydrogenated fluoroethylenes. Furthermore, when high hydrogen/chlorofluoroethane molar ratios are employed, it is possible also to obtain the saturated products resulting from the addition of hydrogen to the fluoro-olefins and chlorofluoro-olefins produced, such as for example CFHCl-CFHCl , $\text{CH}_2\text{F-CH}_2\text{F}$.

According to one preferred embodiment of the present invention, a flow of reagents and hydrogen is continuously fed on the heated catalytic bed.

When leaving the reactor, the vapours are washed with a water solution containing 5-20% by weight of alkaline hydroxide, then they are dried over CaCl_2 and cooled until complete condensation.

For a better understanding of the present invention, the following illustrative examples are given, which, however, are not to be construed as a limitation of the invention.

In the examples, the percentages are by weight, unless otherwise specified.

EXAMPLES 1-10

Into a cylindrical reactor made of steel AISI 316, having an inside diameter of 12 mm and a useful volume of 125 cm³, containing 100 cm³ of activated carbon granules with a palladium content of 0.5% by weight, there was introduced, at atmospheric pressure, a mixture, preheated at 105°C, of hydrogen and difluorotetrachloroethane. Difluorotetrachloroethane consisted of a mixture containing 88% by weight of 1,2-difluorotetrachloroethane and 12% by weight of 1,1-difluorotetrachloroethane. Contact time, reaction temperature and H₂/C₂Cl₄F₂ molar ratio are indicated in Table 1.

When leaving the reactor, the vapours were washed with a 10% NaOH solution, dried and condensed in a trap maintained at -78°C means of dry ice.

On gas-chromatographic analysis, the reaction mixture resulted to be composed of:

- difluoroethylene (C₂H₂F₂)
- chlorodifluoroethylene (C₂HClF₂)
- dichlorodifluoroethylene (C₂Cl₂F₂).

The percentages by weight of the products are reported in Table 1.

The balance to 100 consisted of low boiling by-products. Generally, the unreacted starting product was present only in little amounts.

EXAMPLE 11 (comparative test)

The reaction was carried out according to the modalities of Examples 1-10, with a H₂/reagents molar equal to 5,0, a contact time of 20 seconds and a temperature of 120°C. The following results (% by weight) were obtained:

C₂H₂F₂ 10

C₂Cl₂F₂ 10

The balance to hundred consisted of traces of by-products and of unreacted chlorofluoroethane (78%).

EXAMPLE 12

Into a cylindrical reactor made of steel AISI 316, having an inside diameter of 12 mm and a useful volume of 125 ml, containing 100 ml of activated carbon granules, with a palladium content of 0.5% by weight, thermoregulated at 320°C, there were introduced, at atmospheric pressure,

0.37 moles/h of a mixture, preheated to 105°C, composed of hydrogen and of 1,1-difluorotetrachloroethane in the following molar ratio: H₂/C₂Cl₄F₂ = 2.5.

The vapours leaving the reactor were washed with a NaOH solution at 10%, dried and condensed in a trap maintained at -78°C by means of dry ice.

On gas-chromatographic analysis, the reaction mixture exhibited the following composition:

	(% by weight)
CF ₂ = CCl ₂	15.2
CF ₂ = CHCl	22.3
CF ₂ = CH ₂	37.8
CH ₃ — CHF ₂	9.2

EXAMPLE 13

Into the same reactor and under the same conditions as in Example 12 there were introduced, at atmospheric pressure, 0.37 moles/h of a mixture, preheated to 105°C, of hydrogen and of 1,2-difluorotetrachloroethane in a molar ratio H₂/C₂Cl₄F₂ = 2.5.

The vapours leaving the reactor were washed with a 10% NaOH solution, dried and condensed in a trap maintained at -78°C by means of dry ice. On gas-chromatographic analysis, the reaction mixture exhibited the following composition (% by weight):

CCIF = CCIF 9.8

CHF = CHF 53.2

CCIF = CHF 8.4

CH₂F = CH₂F 8.6

The balance to hundred consisted of low-boiling by-products.

EXAMPLE 14

Into the same reactor and under the same conditions as in Example 12 there was introduced, at atmospheric pressure, a mixture of hydrogen and of 1,1,2-trifluorotrichloroethane in the molar ratio: H₂/C₂Cl₃F₃ = 2,4, said mixture having been preheated to 105°C. The contact time was of 17 seconds. The vapours leaving the reactor were washed with a 10% NaOH solution, dried and con-

densed in a trap maintained at -78°C by means of dry ice. The reaction mixture, subjected to gas-chromatographic analysis, exhibited the following composition:

(% by weight)			
CF_2	=	CHF	30
CF_2	=	CClF	8
$\text{CF}_2\text{H} - \text{CH}_2\text{F}$			35

EXAMPLES 15-17

Under the same conditions as in Example 12, but using as a catalyst Ni on carbon, there was introduced into the usual reactor, at atmospheric pressure, a mixture, preheated to 105°C , composed of hydrogen and of difluorotetrachloroethane (88% by weight of 1,2-difluorotetrachloroethane and 12% by weight of 1,1-difluorotetrachloroethane).

The contact time was of 20 seconds. Three tests were carried out, using different H_2 /difluorotetrachloroethane molar ratios. The vapours leaving the reactor were treated in like manner as in Example 12.

The gas-chromatographic analysis revealed that the reaction products were present in the % by weight reported in Table 2.

The balance to hundred consisted of low-boiling by-products.

EXAMPLE 18

A test was carried out in the same reactor and under the same operative conditions of Examples 15-17, but using Cr on carbon as a catalyst.

The gas-chromatographic analysis revealed that in the product mixture olefin $\text{C}_2\text{Cl}_2\text{F}_2$ was present at 64% by weight.

Table 1

Example	T °C	Molar ratio	Contact time (sec)	$C_2H_2F_2$	C_2HClF_2	$C_2Cl_2F_2$	$C_2H_4F_2$
1	220	2.5	22	51	5.2	10	8.1
2	220	3.6	20	41	3.5	10	24
3	220	1.3	20	13	20	47	2
4	320	2.5	21	55.5	6.5	13.3	6.5
5	320	4.2	20	41	5.0	5.0	25
6	320	1.7	20	11	30	40.0	4.0
7	220	5.0	20	29	2.0	3	43
8	400	2.5	20	61	5.0	2.2	12.3
9	320	3.0	20	51	7	7	11
10	150	2.5	20	15	10	32	3.5

Table 2

Example	Molar ratio	$C_2H_2F_2$	C_2HClF_2	$C_2Cl_2F_2$	$C_2H_2Cl_2F_2$
15	5.0	80	-	-	7
16	2.5	45	37	-	1.5
17	1.3	5	24	62	1.2

Claims

1. A process for preparing fluoroethylenes and chlorofluoroethylenes, which comprises reacting hydrogen with chlorofluoroethanes having at least one chlorine atom on each carbon atom, in gas phase, at a temperature ranging from 150 to 600°C and in the presence of a hydrogenation catalyst.

2. The process according to claim 1, wherein the hydrogenation catalyst is selected from palladium, nickel, chromium, cobalt, platinum, copper and mixtures thereof.

3. The process according to claim 2, wherein the hydrogenation catalyst is selected from palladium and nickel.

4. The process according to any of claims 1 to 3, wherein hydrogen and chlorofluoroethanes are reacted in a molar ratio ranging from 1.0 to 10.

5. The process according to any of claims 1 to 4, wherein the chlorofluoroethanes are selected from 1,2-difluorotetrachloroethane, 1,1-difluorotetrachloroethane, 1,1,2-trichlorotrifluoroethane and mixtures thereof.

6. The process according to any of claims 1 to 5, wherein the reaction temperature ranges from 200 to 400°C.

5

10

15

20

25

30

35

40

45

50

55

6



European Patent
Office

EUROPEAN SEARCH REPORT

Application number

EP 87 11 0386

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.4)
X	EP-A-0 053 657 (ALLIED CORP.) * Claims; page 6, paragraph 1 *	1-4,6	C 07 C 21/18 C 07 C 17/00
X	--- US-A-2 704 775 (J.W. CLARK) * Column 1, line 53 - column 2, line 60 *	1-4,6	
X	--- US-A-2 864 873 (C.B. MILLER et al.) * Claims *	1-2,4-6	
X	--- US-A-3 505 417 (L.E. GARDNER) * Claims *	1-2,4,6	
X	--- US-A-3 636 173 (L.E. GARDNER) * Column 1, line 48 - column 4, line 2 *	1,4-6	TECHNICAL FIELDS SEARCHED (Int. Cl.4)
X	--- CHEMICAL ABSTRACTS, vol. 104, no. 13, 31st March 1982, page 667, abstract no. 109005x, Columbus Ohio, US; & JP-A-60 185 734 (ASAHI GLASS CO., LTD) 21-09-1985 -----	1,5-6	C 07 C 21/00 C 07 C 17/00
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 30-09-1987	Examiner VAN GEYT J.J.A.
CATEGORY OF CITED DOCUMENTS			
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	

EPO Form 1503 03.82

THIS PAGE BLANK (USPTO)

RECEIVED

FEB 04 2005

WELLS ST. JOHN